Mechanism Switch in Mannich-type Reactions. ELF and NCI Topological Analyses of the Reaction between Nitrones and Lithium Enolates

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Abstract: The mechanism of the addition of lithium enolates derived from esters, ketones and aldehydes to nitrones (Mannich-type reaction) has been studied using DFT methods. While the reactions with α -methoxy and α -methyl enolates takes place through a stepwise mechanism, consisting of an initial nucleophilic attack of the enolate to the nitrone carbon followed by a second nucleophilic attack of the nitrone oxygen to the formed carbonyl group, the reaction with α -unsubstituted enolates takes place through a one-step mechanism. The IRC analysis shows the presence of a hidden intermediate in agreement with one kinetic step two stages process. The topological analysis of the electronic localization function (ELF) confirms that only when the first C-C bond is formed, does the C-O bond formation begin. The NCI analyses, are also in agreement with the formation of intermediates for α -methoxy and α -methyl enolates and a highly asynchronous one-step process in the case of α -unsubstituted enolates.

Introduction

Mannich-type reactions are probably the most popular approach for the synthesis of β -amino carbonyl compounds.^[1] The direct addition of enolates to a variety of functionalities including imines^[2] as well as other C=N groups such as nitrones, hydrazones and iminium salts^[3] is especially useful to create different types of β -nitrogenated carbonyl derivatives in a single synthetic operation. In particular, the use of nitrones as substrates has received considerable attention because they lead to β -aminocarbonyl functionalities in which the nitrogen group is at an intermediate oxidation state (Scheme 1). The final product can be a β -hydroxyamino carbonyl derivative or the corresponding isoxazolidin-2-one obtained after an intramolecular cyclization. In addition, the Mannich-type reaction of nitrones has provided access to enantiomerically pure compounds of biological and pharmacological interest including aminosugars,^[4] iminosugars,^[5] nucleoside analogues,^[6]

sphingosines^[7] and aminoacids.^[8]

Scheme 1. Mannich-type reactions of nitrones

Several nucleophiles can be used in the reaction,^[9] the most commonly employed being enolates derived from esters in the form of lithium,^[6a] sodium,^[6a] boron^[8b,10] and titanium salts.^[8b,10-11] Silyl enolates derived from esters (silyl ketene acetals) have also extensively used in the presence of Lewis acids.^[4b,6b,c 8c,e,12] Recently, enol silanes formed *in situ*, derived from ketones, amides and thioesters have been reported to add to nitrones in the presence of trialkylsilyl trifluoromethane-sulfonates.^[13] On the contrary, there are only two examples regarding the reaction between nitrones and metal enolates derived from ketones^[14] and a self-catalyzed Mannich-type reaction between nitrones and 1,3-dicarbonyl compounds (without any base) has been recently reported.^[15] To the best of our knowledge, no previous reports on the reaction with metal enolates derived from aldehydes are documented.

From a mechanistic point of view, the addition of silyl ketene acetals to nitrones presented some controversy since the reaction was initially postulated to take place through a stepwise mechanism^[8c,16] whereas a concerted mechanism involving a pentacoordinated silicon was also invoked on the basis of semiempirical calculations.^[8e] On the other hand, a different semiempirical study pointed out that the mechanism of the reaction could change from concerted to stepwise depending on the Lewis acid used as activating agent.^[17] Further DFT calculations with very simple models in gas phase and without considering Lewis acids (which are required for the advance of the reaction) presented the reaction like a typical concerted inverse-demand 1,3-dipolar cycloaddition.^[18] Finally, consideration of a more realistic scenario taking into account both the presence of Lewis acids and solvent effects confirmed that both concerted and stepwise mechanisms are competitive.^[19]

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Experimental and theoretical investigations have demonstrated that the reactions of organometallic reagents, such as organolithium^[19-20] and Grignard^[21] derivatives, with nitrones take place through the initial formation of a complex **SC** (Scheme 2). When the nucleophile is a lithium enolate (usually derived from an ester) the final product of the reaction is an isoxazolidine. This heterocycle could be formed from the initial complex **SC** through either a concerted mechanism (Scheme 2, A) or a typical nucleophilic addition stepwise mechanism (Scheme 2, B). During past investigations we confirmed the stepwise mechanism for α -methoxyenolates derived from esters ($Z=OR$).^[19,20c] In that case, the stepwise mechanism is favored because of the stabilization of the developing positive charge in **TS-B1** by the $Z=OR$ group. However, it remains unstudied the case of lithium enolates derived from aldehydes and ketones (Scheme 2, $Z = H, R$) in which such stabilization is not present and the concerted mechanism could be an option.

Scheme 2. One-step (A) and stepwise (B) mechanisms for the addition of lithium enolates to nitrones.

Despite the apparent similarity between lithium enolates derived from aldehydes, ketones and esters all these species are rather different if we consider their electronic properties,^[22] which are crucial for the stability of postulated intermediates. In order to shed some light on the preferred pathways (concerted or stepwise) it is necessary a full theoretical study considering in detail all the possible paths. Herein, we report a DFT study on the reaction of lithium enolates derived from esters, ketones aldehydes. The study includes a detailed analysis of all points of the intrinsic reaction coordinates (IRCs) by using NCI and ELF topological analyses that will allow discerning the concertedness of each process.

Computational Methods

All of the calculations were performed using the Gaussian09 program.^[23] Molecular geometries were optimized with the M06-2X functional^[24] in conjunction with cc-pVTZ basis set.^[25] Truhlar and co-workers reported that for examining barrier heights, a minimally augmented basis set like the Dunning cc-pVTZ is appropriate.^[26] It was not necessary to augment the pTZV basis with extra diffuse functions, as tests carried out by using aug-cc-pVTZ basis, resulted in changes in the relative energies of less than 1 kcal/mol while making calculations considerably more time-consuming. Moreover, the use of M06-2X in conjunction with cc-pVTZ basis has provided excellent results in related calculations with nitrones.^[27] We can therefore expect that the computed values should be sufficiently reliable to be able to draw meaningful conclusions. Analytical second derivatives of the energy were calculated to classify the nature of every stationary point, to determine the harmonic vibrational frequencies, and to provide zero-point vibrational energy corrections. The thermal and entropic contributions to the free energies were also obtained from the vibrational frequency calculations, using the unscaled frequencies. All transition structures were characterized by one imaginary frequency and were confirmed to connect to reactants and products by intrinsic reaction coordinate (IRC) calculations.^[28] The IRC paths were traced using the second order González-Schlegel integration method.^[29] Solvent effects were calculated using the continuum solvation model (PCM)^[30] using a dielectric constant of 7.4257 to simulate THF, and microsolvation of the lithium atom was considered by explicit inclusion of dimethyl ether ligands (to reduce computational cost we replace discrete THF molecules by dimethyl ether units) to complete the coordination sphere of lithium, according to previous reports by Domingo and co-workers.^[31] A four-fold (tetrahedral) coordination sphere was chosen for the lithium atom in agreement with previous studies.^[32] NCI (non-covalent interactions) were computed using the methodology previously described.^[33] Data were obtained with the NCIPLOT program.^[34] A density cutoff of $\rho=0.1$ a.u. was applied and the pictures were created for an isosurface value of $s=0.4$ and colored in the $[-0.02, 0.02]$ a.u. $\text{sign}(\lambda_2)\rho$ range using VMD software.^[35] The electronic structures of stationary points were analyzed by the topological analysis of the gradient field of electron localization function (ELF)^[36] developed by Silvi and Savin.^[37] The ELF study was performed with the TopMod program^[38] using the corresponding monodeterminantal wavefunctions of the all structures of the IRC. The topological analysis of the gradient field of ELF has showed to be a powerful tool for the study of the bonding changes along an organic reaction.^[39] Advantages and drawbacks of NCI index, compared with ELF^[40] and QTAIM (quantum theory of atoms in molecules)^[41] have been reported. Structural representations were generated using CYLView.^[42] Animation given in the supporting material was created by extracting and processing all points of the IRC with an in-house program and saving the corresponding images to create an animated GIF. The lithium enolate derived from methyl acetate (**ENa**, $Z = OMe$), the actual reagent employed in previous experimental reports,^[20c, 43] was

used in the calculations. For the purpose of comparison and in order to study the only differences exerted by diverse groups vicinal to the carbonyl function, the corresponding enolates derived from acetone (**ENb**, Z = Me) and acetaldehyde (**ENc**, Z = H) were chosen as models. To reduce computational cost, the model nitron **NI** maintaining the fundamental characters of a real N-substituted (Z)-nitron (the preferred configuration for aldolones), has been chosen.

Results and Discussion

Mechanistic study. Even though a lithium enolate can be expressed as ROLi , its composition in a real solution is far more complex. In polar solvents like THF (and in the absence of chelating agents) lithium enolates are cubic tetramers,^[44] although in some cases dimers can be formed and, in the presence of additives, monomers can also be present.^[45] A computational study carried out with the lithium enolate derived from acetaldehyde demonstrated that monomeric species is important in the equilibrium due to its high solvation energies,^[46] although more recent calculations showed that tetramer is the major species in THF, monomeric species being preferred in the presence of chelating agents.^[47] Additional computational studies reported that modeling by coordination to dimethyl ether and dielectric solvation reduces considerably the exothermicity of the aggregation.^[48] An experimental study with the lithium enolate of α -phenylcyclohexanone demonstrated that an equilibrium exists between the monomer and the dimer, the former being more reactive in alkylation reactions.^[49] Thus, it is reasonable to assume that, with independence of the structural type of the lithium enolate in solution, the nitron can break the aggregates (in a similar way to the reactions with Grignard reagents^[50]) and form an initial complex **SC** as considered for other reactions.^[48]

We consider six different approaches, leading to the corresponding transition structures, between nitron **NI** and monomeric enolates **ENa**, **ENb** and **ENc**, corresponding to three staggered orientations and two different faces of the enolate (Scheme 3). From these six approaches only three of them, leading to **a**, **b**, and **c** series allow the preferred coordination of both reagents to the lithium atom. Thus, for those leading to **d**, **e**, and **f** series, corresponding to a direct approach without formation of an initial complex, an additional molecule of solvent has been added to complete a fourfold coordination sphere for lithium.^[51]

We first revisited the mechanism of the nucleophilic addition of α -methoxyenolate **ENa** to nitron **NI**. In our previous report^[19] we considered **IN** (Scheme 2), formed from complex **SC** through **TS-B1**, as the final product of the reaction. However, the reaction can continue through two alternative diastereomeric channels to form cyclic *cis* and *trans* products **PR** which evolve to the isoxazolidin-2-one **4** observed experimentally (Scheme 4).

[6a,20c,52]

Scheme 3. Approaches between nitron **NI** and enolates **ENa-c**.

Formally, the reaction between **NI** and **ENa** to form **P1a** and **P1b** can be considered a [3+2] cycloaddition. The analysis of the potential energy surface showed the direct approach higher in energy and thus the corresponding transition structures **TS1d**, **TS1e** and **TS1f** were not further considered (Figure 1). On the other hand, formation of a complex **C1** resulted in a stabilization of 4.2 kcal/mol. Starting from **C1** the three transition states **TS1a**, **TS1b** and **TS1c**, corresponding to the formation of a C-C bond between the most nucleophile center of the enolate **ENa** (the unsubstituted methylene) and the most electrophilic center of the nitron **NI** (the azomethine carbon) through the two faces of the enolate, were located. The complete mechanism is given in Scheme 4 while the energy profile for the reaction and main geometrical features of stationary points corresponding to the formation of **P1a** and **P1b** are given in Figure 1.

The IRC analyses confirmed **C1** as the starting point for the three transition structures, and **IN1b** as the final point for both **TS1b** and **TS1c**; for **TS1a** in which the addition of the nitron takes place by the other face of the enolate **IN1a** was identified as the final point. The most stable transition state corresponds to **TS1a** with an energy barrier of 6.4 kcal/mol whereas **TS1b** and **TS1c** present barriers of 7.2 and 8.5 kcal/mol, respectively. According to a classical Boltzmann distribution analysis, **TS1a** accounts for 80% of all the transition structures but in terms of the mechanism it is irrelevant because all the approaches are stepwise.

Scheme 4. Reaction between nitron **NI** and α -methoxy enolate **ENa**

Figure 1. Energy diagram (M06-2X/cc-pVTZ/PCM=THF) and stationary points for the reaction between nitron **NI** and α -methoxy enolate **ENa**. Relative free energy values (ΔG_{298}) are given in kcal/mol.

The distances of the forming bond in **TS1a**, **TS1b** and **TS1c** are 2.18, 2.14 and 2.15 Å, respectively, whereas the distances between the nitron oxygen and the enolate carbon linked to the two oxygen atoms are 3.43, 3.11 and 3.54 Å for **TS1a**, **TS1b** and **TS1c**, respectively thus confirming the absence of a close interaction between the two reactive centers. Consequently, the reaction is a typical nucleophilic addition. Indeed, any attempt of locating concerted transition structures in which the formation of

the two bonds could take place in a concerted (although asynchronous) manner, failed. The stability of **IN1a** and **IN1b** was confirmed after the corresponding optimizations which showed them to be 11.8 and 13.2 kcal/mol below the ground state, respectively. Both intermediates could also interconvert through a process of decooordination of the lithium atom and rotation of the ester moiety. The formation of **IN1a** and **IN1b** is followed by the intramolecular attack of the hydroxyamino group

to the ester carbonyl, via the planar four-membered ring transition states **TS4a** and **TS4b**, leading eventually to the lithium coordinated orthoesters **P1a** and **P1b**, respectively. The forming C-O bond lengths in **TS4a** and **TS4b** are 2.06 Å and 1.94 Å, respectively and in both cases the IRC analysis showed **IN1a** and **IN1b** as the starting points. Also, in both **TS4a** and **TS4b** the forming isoxazolidine ring adopts an envelope conformation. The *channel b* is favored over the *channel a* since the nucleophilic attack of the oxygen atom to the ester carbonyl in the latter is found to be **sterically** better positioned.

The whole processes are exergonic by 10.6 kcal/mol for *channel a* and 14.5 kcal/mol for *channel b*. As a result, whereas *channel a* is kinetically preferred because **TS1a**, at the rate-determining step, is the most stable, *channel b* is more favorable thermodynamically. However, the selectivity cannot be experimentally observed because after quenching the reaction the corresponding orthoesters eliminate the methoxy group to give isoxazolidine-2-one **4**. We have not studied computationally the final step leading to **4** because it has no relevance in the mechanism of the addition reaction. Thus, in the case of α -methoxyenolates the stability of intermediates **IN1a** and **IN1b** is ultimately responsible for the process to be stepwise. Indeed, in some similar cases it has been observed experimentally^[53] the obtention of free hydroxylamine directly derived from those complexes.

In the case of ketone enolate **ENb**, calculations also establish that the direct approach, in which only **ENb** is coordinated to lithium, are prohibitively high in energy, with energy barriers of 30.2, 24.4 and 29.8 kcal/mol for **TS2d**, **TS2e** and **TS2f**, respectively. On the other hand, starting from complex **C2**, located at 4.2 kcal/mol below the ground state, energy barriers of 8.3, 9.9 and 11.7 kcal/mol were found for

TS2a, **TS2c** and **TS2b**, respectively (Scheme 5, Figure 2). The geometrical features as well as the energy profile are given in Figure 2.

The IRC analysis for **TS2a** and **TS2c** (both attacking by the same face of the enolate) confirmed intermediates **IN2a** and **IN2b** as the final points of the reaction. However, the same analysis for **TS2b** (corresponding to the attack for a different face of the enolate) indicated the process as concerted, the product **P2a** being the final point. The geometries of transition structures **TS2a-c** (R = Me, Scheme 5) are similar to the corresponding partners **TS1a-c** (Scheme 4) but the electronic features are different. In the case of **TS1b**, the presence of the methoxy group contributes to stabilize the developing partial positive charge at the carbonyl carbon atom; the C-O distance of 3.11 Å indicates that electrostatic interaction is not enough for causing the collapse of the second forming bond. Under these circumstances the intermediate **IN1b** is enough stable to exist. On the other hand, in **TS2b** the methyl group is not capable of stabilizing the above mentioned partial positive charge and, consequently, the electrostatic interaction between carbon and oxygen atoms is stronger (as revealed by a shorter C-O distance of 3.05 Å). In this scenario the C-O interaction collapses to a bond and the reaction takes place in one single kinetic step. **This sort of spontaneous downhill process is well known and it has also been observed by other authors.**^[54] The *endo* orientation of the methyl group causes unfavorable steric interactions that explain the higher energy of **TS2b** with respect to **TS2a** and **TS2c**. As in the case of α -methoxyenolate, the formation of **IN2a** and **IN2b** is followed by an intramolecular attack through **TS5a** and **TS5b** leading to orthoesters **P2a** and **P2b**, respectively (Scheme 5, R = Me).

Scheme 5. Reaction between nitrone **NI** and enolates **ENb,c**

Figure 2. Energy diagram (M06-2X/cc-pVTZ/PCM=THF) and stationary points for the reaction between nitron **NI** and α-methoxy enolate **ENb**. Relative free energy values (ΔG_{298}) are given in kcal/mol.

Notably, the IRC analysis of the concerted pathway showed a shoulder revealing the presence of a so-called hidden intermediate (Figure 3).^[55] According to this analysis and the excessively long forming C-O bond (3.05 Å), only when the transition state is passed and the C-C bond is formed, does the formation of the second C-O bond start. This process consisting in two consecutive chemical events (formation of C-C and C-O bonds) is in agreement with a typical *one-step-two-stages* reaction according to Domingo and co-workers^[39b] and similar to that observed for the reaction between nitrones and lithium ynolates.^[27b] Indeed, on going from **TS2b** to cycloadduct **P2b**, through IRC, intermediate structures (see below the ELF analysis) have the C-C bond already formed whereas the C-O bond formation is very delayed. However, steric reasons due to the *inside* orientation of the methyl group make **TS2b** higher in energy and we can conclude that this path is not preferred for this reaction. The Boltzmann distribution analysis predicted that **TS2a**, corresponding to the stepwise process, accounts for almost 95% of all the transition structures while **TS2b**, corresponding to the one-step process, accounts for only 5%. Consequently, despite the presence of the concerted path, the

reaction between nitron **NI** and α-methyl enolate **ENb** takes place through a stepwise mechanism with a barrier of 8.3 kcal/mol at the rate-limiting step.

Figure 3. Computed (M06-2X/cc-pVTZ/PCM=THF) backwards intrinsic reaction coordinate (IRC) for the reaction between nitron **NI** and enolate **ENb** showing the relative energy (top) and the gradient norm showing a prominent hidden intermediate (bottom).

A similar situation accounts in the reaction between nitron **NI** and enolate **ENC**. Again, the three transition structures corresponding to the direct approach present very high energy barriers (30.8, 27.9 and 29.2 kcal/mol for **TS3d**, **TS3e** and **TS3f**, respectively). The free energy barriers for lithium-coordinated transition structures **TS3a**, **TS3b** and **TS3c** are calculated to be 12.3, 10.5 and 12.1 kcal/mol, respectively. The geometrical features as well as the energy profile are given in Figure 4.

As observed for α-methyl enolate, **TS3b** appeared as a highly asynchronous transition structure with a long C-O forming bond (2.99 Å) in comparison with the C-C forming bond (2.10 Å). The IRC calculation confirmed the concertedness of the reaction revealing no intermediates between **TS3b** and **P3b**. However, contrary to α-methyl enolate, transition state **TS3b**,

corresponding to the one-step process, showed to be the most stable (by 1.6 kcal/mol) confirming a change of mechanism from enolate **ENb** to enolate **ENc**. In fact, the Boltzmann distribution analysis indicated in this case that **TS3b** accounts for almost 92% of all the transition structures while **TS3a** and **TS3c**, corresponding to stepwise processes, account for ca. 3% and 5%, respectively. The steric contact of the *N*-methyl group is more unfavorable for the methyl group (**TS2b**) than for the hydrogen atom (**TS3b**), thus predicting a lower barrier for the α -unsubstituted enolate **ENc** (Figure 4). The unfavorable formation of **IN3a** and **IN3b** is followed by an intramolecular attack through **TS6a** and **TS6b** leading to orthoesters **P3a** and **P3b**, respectively (Scheme 5, R = H). The IRC calculation for the concerted process show similar features to those observed in the case of α -methyl enolate for **TS2b**. It confirms that **C3** and **P3a** are connected by **TS3b** without intermediates but a hidden intermediate is also present (Figure 5). Thus, in the case of the α -unsubstituted enolate **ENc**, the reaction takes place preferentially along a concerted two-stage one-step mechanism with an energy barrier of 10.5 kcal/mol.

Figure 4. Energy diagram (M06-2X/cc-pVTZ/PCM=THF) and stationary points for the reaction between nitron **NI** and α -methoxy enolate **ENc**. Relative free energy values (ΔG_{298}) are given in kcal/mol.

Figure 5. Computed (M06-2X/cc-pVTZ/PCM=THF) intrinsic reaction coordinate (IRC) for the reaction between nitron **NI** and enolate **ENc** showing the relative energy (top) and the gradient norm showing a prominent hidden intermediate (bottom).

NCI and ELF analyses. The topological analysis of ELF has recently demonstrated to be of great utility in analyzing C-C bond formation in a variety of non-polar, polar and ionic organic reactions.^[56] The NCI analysis^[33] has also demonstrated their utility in the analysis of several reactions^[55b,57] including nucleophilic additions to C=N bonds.^[58] We have carried out the complete ELF and NCI analyses for the IRCs corresponding to the most stable paths of the addition reactions of enolates **ENa-c** to nitron **NI** (For animations showing movies of the reactions illustrating both ELF and NCI analyses see supporting material). The numbering used for the analyses is illustrated in Figure 6. ELF basin populations of selected points on the IRC including initial and final points, transition structures, intermediates and points indicating bond formation are given in the supporting information.

Figure 6. Numbering used for ELF and NCI analyses.

In the case of the reaction between **NI** and **ENa** leading to **TS1**, the double C3=N1 bond is transformed into the single C3-N1 bond, the double C6=C7 bond is transformed into the single C6-C7 bond and the C3-C7 bond is created. The ELF descriptors corresponding to this step are presented in Figure 7. The two disynaptic basins associated to the C3=N1 and C6=C7 double bonds are merged, in **TS1a** (point 58 of the IRC), each other to become one indicating the transformation of the double bonds into single one; in fact the electronic populations decreased slightly for both C3-N1 and C6-C7 bonds. The decreasing of the electronic population of C3-N1 and C6-C7 bonds continues during C3-C7 bond formation (points 55 and 54 of the first IRC) and simultaneously a monosynaptic basin appeared at C7. At point 55 ($d(\text{C3}, \text{C7}) = 2.04 \text{ \AA}$; $d(\text{C6}, \text{O2}) = 3.45 \text{ \AA}$) two monosynaptic basins, $V(\text{C3})$ and $V(\text{C7})$, appeared at the reacting centers. These basins are associated to the two centers responsible for the subsequent bond formation. Indeed, at the next point on the IRC (point 54) they have merged into a new disynaptic basin, $V(\text{C3}, \text{C7})$ confirming the C3-C7 bond formation.

Figure 7. Most relevant ELF attractors at selected points of the backwards IRCs of the stepwise reaction between nitrone **NI** and enolate **ENa**.

In the second step of the reaction, at **TS4a** (point 28 of the IRC) no monosynaptic basins appeared at the centers

responsible of the formation of the second bond, C6 and O2. It is at point 21 when two monosynaptic basins, $V(\text{C6})$ and $V(\text{O2})$, appeared. At **P20** they have merged into a disynaptic one confirming the formation of the new bond

The NCI analysis (Figure 8) for this reaction corroborates and complements the data observed during the ELF analysis. The starting complex, **C1**, shows a typical attractive interaction (green surface) between the π systems corresponding to electron-rich enolate C=C bond and the electron-poor nitron C=N bond. At the intermediate **IN1a** a clear non-covalent interaction is observed between C6 and O2 (blue surface) in agreement with a relatively short distance of $2.70 \text{ \AA}$ and despite the formation of the second C6-O2 bond does not have started, as mentioned above. This observation is in agreement with that made for anionic stepwise [3+2] cycloadditions by Schelyer and co-workers who considered this sort of interaction as strictly electrostatic.^[59] More recently, we have also observed the same type of interactions in the stepwise cycloaddition between nitron ylides and alkenes.^[20b] For transition structures **TS1a** and **TS4a** the incipient formation of the new bonds is evidenced by the typical toroidal blue surfaces (Figure 8). At **TS1a** is evident that the interaction between C6 and O2 is negligible only being appreciable at the following stationary point **IN1a**.

Figure 8. NCI analysis of relevant points **C1**, **IN1a**, **TS1a** and **TS4a** corresponding to the stepwise reaction between nitrone **NI** and enolate **ENa**.

For the concerted reaction between nitrone **NI** and enolate **ENc** the attractor positions for points indicating bond formation are illustrated in Figure 9. For this reaction the ELF analysis of the attractors for **C3** shows, in a similar way to **C1**, two disynaptic basins associated each one to the C3=N1 and C6=C7 double bonds of the nitrone and enolate moieties, respectively. At **TS3b** (point 60 of the IRC) the C6-C7 and C3-N1 bonding regions are characterized by $V_1(\text{C6}, \text{C7})$ and $V_1(\text{C3}, \text{N1})$ disynaptic basins, which showed loss of electron density associated to the creation of the new C3-C7 bond. A monosynaptic basin $V(\text{C7})$ is observed and at **P62** a new monosynaptic basin, $V(\text{C3})$, appears

and the electron density of $V(C7)$ increases. At the following point, **P63**, the two monosynaptic basins $V(C3)$ and $V(C7)$ have merged into a new disynaptic basin $V(C3,C7)$. Notably, **P62** and **P63** associated with the first stage of the concerted process have similar electronic structures to **P55** and **P54** associated with the stepwise addition of **ENa** (see above). This indicates a similar arrangement in the formation of the first C-C bond, independently of the appearance of a further intermediate, whose stability (or existence) depends on electronic features that could stabilize such stationary point. Indeed, **P63** resembles geometrically **IN1a** and the absence of $V(C6,O2)$ attractor confirms that formation of the second bond has not begun. At **P96** both C3,C7 distance (1.54 Å) and the presence of $V(C3,C7)$ attractor indicate the complete formation of the C3-C7 bond. At the same time, two monosynaptic basins, $V(C6)$ and $V(O2)$ appear. These basins merge, at **P97**, into a new disynaptic basin ($V(C6,O2)$) responsible of the formation of the second bond.

Figure 9. Most relevant ELF attractors at selected points of the IRCs of the concerted reaction between nitron **NI** and enolate **ENc**.

The NCI analysis of the starting complex **C3**, the transition structure **TS3b** and the two points, **P63** and **P97**, in which the formation of the bonds has just taken place is illustrated in Figure 10. Similarly to **C1**, complex **C3** shows an attractive interaction (green surface) between the π systems corresponding to electron-rich enolate C=C bond and the electron-poor nitron C=N bond. **TS3b** shows the interaction corresponding to the forming bond (toroidal blue surface). **TS3b** and **P63** are very similar in their geometrical structure and both show a slight attractive interaction between C6 and O2 in a similar way (although weaker) to that observed for **IN1** in the stepwise addition of **ENa**. Although an intermediate is not formed it is evident that the second bond (C6-O2) is not formed at this stage of the reaction, being completely formed only at **P97**.

The case of the reaction between nitron **NI** and α -methyl enolate **ENb** can be considered as an intermediate situation which, however, opts by a stepwise mechanism due to

unfavourable steric interactions still present between the methyl group of the α -enolate and the *N*-methyl group. Both the ELF and NCI analyses are rather similar to those discussed above for the reaction with enolate **ENa** by just replacing the α -methoxy group by the α -methyl group. The same applies for the non-preferred concerted path which is rather similar to those found for enolate **ENc** by just replacing the α -methyl group by an hydrogen atom (For the complete analyses of both concerted and stepwise pathways of the reaction between nitron **NI** and enolate **ENb** see supporting information).

Figure 10. NCI analysis of relevant points **C3**, **TS3a**, **P63** and **P97** corresponding to the concerted reaction between nitron **NI** and enolate **ENc**.

Conclusions

The addition of lithium enolates to nitrones takes place through the initial coordination of the nitron to the lithium atom. Then, the intramolecular attack of the enolate moiety to the nitron from initial complexes can take place by two different faces of the enolate. This causes that the α -substituent of the enolate (OMe, Me or H) can adopt *inside* and *outside* orientations with respect to the nitron (Figure 11).

Figure 11. Preferred approaches and change of mechanism for enolates derived from esters and ketones ($Z = \text{OMe, Me}$), and aldehydes ($Z = \text{H}$). (dashed lines indicate forming bonds).

The *inside* position is sterically more demanding because of unfavorable interactions with the *N*-methyl group of the nitron and thus, the *outside* orientation is preferred. However, this approach involves a large separation between C and O atoms responsible of the formation of the second C-O bond. This excessively long C-O distance, minimizing attractive electrostatic interactions, avoids bonding between the interacting orbitals at the TS, and when an electron-donor substituent (OMe) is present the reaction is stepwise because of the electronic stabilization of the intermediate (addition of **ENa**). In the absence of α -substituent the *inside* orientation is preferred (shortening the C-O distance) and the reaction change its mechanism to concerted through a highly asynchronous transition structure (addition of **ENc**). The reaction with the α -methyl enolate (**ENb**) represents an intermediate situation in which there is a substituent stabilizing in less extent the intermediate but still causing unfavorable steric interactions. Consequently, the concerted path appeared but is not the preferred one and the stepwise pathway shows a higher barrier (8.3 kcal/mol) than in the case of α -methoxy enolate (6.4 kcal/mol).

The one-step processes are so asynchronous that they are more in agreement with a reaction which takes place in one single kinetic step but in two stages. This concept has been introduced by Domingo and co-workers^[60] and it is evidenced from the ELF analyses of the corresponding IRC calculations. From these analyses of the changes of bonding along the reaction coordinate we can conclude that, in the one-step processes, the formation of the second C-O bond only begins when the first C-C bond is completely formed. Consequently, these concerted highly asynchronous reactions do not follow a typical cyclic electron-reorganization as supported by the presence of hidden intermediates in the corresponding IRCs. As predicted by Rzepa and co-workers,^[55b] stereoelectronic influence on the geometry induces the system to form a real intermediate.

In summary, while the reaction with α -unsusbtituted enolate **ENc** takes place along a one-step two-stage mechanism, the presence of a substituent at the α -position in enolate (**ENa** and **ENb**) able to both lengthen the C-O distance -avoiding orbital interactions- and stabilize the corresponding intermediate, switches the mechanism to a stepwise process.

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Depending on the type of enolate the reaction mechanism changes from one-step to stepwise